



Annex 1

Claims

- 1. Use of a polyurethane in the manufacturing of a composition or kit used for filling or short-circuiting vascular cavities, where the said polyurethane used can be dissolved is soluble in a solvent or a solvent mixture mingling with body fluids and the polyurethane is the main component responsible for binding into the vascular cavities.
- 2. The use according to claim 1, where a composition is manufactured, which contains the polyurethane dissolved in a solvent or a solvent mixture mingling with body fluids, optionally combined with usual auxiliaries.
- 3. The use according to claim 1, where a kit is manufactured, which contains the following components:
- a) a polyurethane which can be dissolved is soluble in a solvent or a solvent mixture mingling with body fluids, optionally together with other usual auxiliaries,
- b) one or more solvents or solvent mixtures mingling with body fluids, in which the said polyurethane can be dissolved is soluble, optionally together with other usual auxiliaries,
- c) optionally other usual auxiliaries.,
 where the components given above are formulated separately or some of them are formulated in
 a common subunit.
- 4. The use according to any of claims 1 to 3, where the solvent is DMSO or EtOH or their mixture, preferably the mixture of them in the a volume ratio of 1:10 10:1, more preferably of 1:3 3:1.
- 5. The use according to any of claims 1 to 4, where the main diol component of the polyurethane is characterized by the general formula of $HO-R^1-OH$, where R^1 stands for a C_1-C_8 alkylene group, preferably 1,4-buthanediol.
- 6. The use according to claim 5, where 50 to 95 % of the main diol component is in polyether form, preferably in polytetrahydrofurane form.
- 7. The use according to claims 1 to 6, where the main diisocyanate component of the polyurethane is one or more compound(s) selected from the following ones: group consisting of 2,4- or 2,6- toluylene-diisocyanate (TDI), 1,6-hexane-diisocyanate and diphenyl-methane-4,4'-diisocyanate is preferred.





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- 8. The use according to claim any of claims 1 to 7, where the composition or kit contains a polyurethane solution, usable for filling aneurisms, having a viscosity higher than 150 mPa.s, preferably higher than 250 mPa.s at 23 °C.
- 9. The use according to any of claims 1 to 7, where the composition or kit contains a polyurethane solution, usable for filling angiomas and vascularized tumors, having a viscosity is lower than 1 000 mPa.s, preferably lower than 250 mPa.s at 23 °C.
- 10. The use according to any of claims 1 to 9, where the molecular mass of the polyurethane used is 4 000 to 70, 000 Dalton, preferably 20, 000 to 35, 000 Dalton.
- 11. The use according to any of claims 1 to 10, where the composition or kit contains contrast material as an auxiliary, preferably selected from the following group: a substance containing tantalum, iodine, barium, tungsten and/or bismuth, of which micronized tantalum powder, tantalum-oxide, barium-sulphate and tungsten are more preferred.
 - 12. Composition for filling or short-circuiting vascular cavities, containing
- a polyurethane which can be dissolved ssoluble in a solvent or a solvent mixture mingling with body fluids and which is the main component responsible for binding into the vascular cavities.
 - optionally dissolved in a solvent or a solvent mixture mingling with body fluids, and optionally combined with usual auxiliaries.
- 13. A composition according to claim 12, which contains the polyurethane dissolved in a solvent or a solvent mixture mingling with body fluids, optionally combined with usual auxiliaries.
- 14. A therapeutic product containing the components formulated separately (kits of parts), which can be used for filling or short-circuiting vascular cavities and contains the following components:
- a) a polyurethane that can be solvedwhich is soluble in a solvent or a solvent mixture mingling with body fluids and which is the main component responsible for binding into the vascular cavities, optionally together with other usual auxiliaries,
- b) one or more solvents or solvent mixtures mingling with body fluids, in which the said polyurethane can be dissolved is soluble, optionally together with other usual auxiliaries,
 - c) optionally other usual auxiliaries.,





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where the components given above are formulated separately or some of them are formulated in a common subunit.

- 15. A composition or a therapeutic product according to any of claims 12 to 14, where the solvent is DMSO or EtOH or their mixture, preferably a mixture of them in the volume ratio of 1:10 10:1, more preferably of 1:3 3:1.
- 16. A composition or a therapeutic product according to any of claims 12 to 15, where the main diol component of the polyurethane is characterized by the general formula of $HO-R^1$ -OH, where R^1 stands for a C_1-C_8 alkylene group, preferably 1,4-buthanediol.
- 17. A composition or a therapeutic product according to claim 16, where 50 to 95 % of the main diol component is in polyether form, preferably in polytetrahydrofurane form.
- 18. A composition or a therapeutic product according to any of claims 12 to 17, where the main disocyanate component of the polyurethane is one or more compound(s) selected from the following ones:group consisting of 2,4- or 2,6- toluylene-diisocyanate (TDI), 1,6-hexane-diisocyanate and diphenyl-methane-4,4'-diisocyanate (MDI), of which diphenyl-methane-4,4'-diisocyanate is preferred.
- 19. A composition or a therapeutic product according to any of claims 12 to 18, where the composition or therapeutic product contains a polyurethane solution, usable for filling aneurisms, having a viscosity higher than 150 mPa.s, preferably 250 mPa.s at 23 °C.
- 20. A composition or a therapeutic product according to any of claims 12 to 18, where the composition or therapeutic product contains a polyurethane solution, usable for filling angiomas and vascularized tumors, having a viscosity lower than 1 000 mPa.s, preferably lower than 250 mPa.s at 23 °C.
- 21. A composition or a therapeutic product according to any of claims 12 to 20, where the molecular mass of the polyurethane used is 4 000 to 70, 000 Dalton, preferably 20, 000 to 35, 000 Dalton.
- 22. A composition or a therapeutic product according to any of claims 12 to 21, where the composition or therapeutic product contains a contrast material as an auxiliary, preferably selected from the following group: a substance containing tantalum, iodine, barium, tungsten or bismuth, of which micronized tantalum powder, tantalum-oxide, barium-sulphate and tungsten are more preferred.





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- 23. Use of a polyurethane for filling vascular cavities, where the applied polyurethane is dissolved in a solvent or solvent mixture mingling with body fluids, and optionally combined with usual auxiliaries.
- 24. The use according to claim 23, where the solvent is either DMSO or EtOH or their mixture, preferably a mixture of them in the volume ratio of 1:10 10:1, more preferably of 1:3 3:1.
- 25. The use according to any of claims 23 and or 24, where the main diol component of the polyurethane is a diol characterized by the general formula of HO-R¹-OH, where R¹ stands for a C₁-C₈ alkanediyl group, preferably 1,4-buthanediol.
- 26. The use according to claim 25, where 50 to 95 % of the main diol component is in polyether form, preferably in polytetrahydrofurane form.
- 27. The use according to any of claims 23 to 26, where the main disocyanate component of the polyurethane is one or more compound(s) selected from the following ones: group consisting of 2,4- or 2,6-toluylene-diisocyanate (TDI), 1,6-hexane-diisocyanate and diphenyl-methane-4,4'-diisocyanate (MDI), of which diphenyl-methane-4,4'-diisocyanate is preferred.
- 28. The use according to any of claims 23 to 27, where a polyurethane solution applicable for filling aneurisms is used having a viscosity higher than 150 mPa.s, preferably 250 mPa.s at 23 °C.
- 29. The use according to any of claims 23 to 27, where a polyurethane solution applicable for filling angiomas and vascularized tumors is used having a viscosity lower than 1 000 mPa.s, preferably lower than 250 mPa.s at 23 °C.
- 30. The use according to any of claims 23 to 29, where molecular mass of the polyurethane used is 4 000 70, 000 Dalton, preferably 20, 000 35, 000 Dalton.
- 31. The use according to any of claims 23- to 30, where the contrast material is used as an auxiliary, preferably selected from the following group: a substance containing tantalum, iodine, barium, tungsten or bismuth, of which micronized tantalum powder, tantalum-oxide, barium-sulphate and tungsten are more preferred.